## IN THE CLAIMS:

## Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims**

- 1-9. (Cancelled)
- 20. (Currently Amended): A method of treating <u>arthritis</u> an autoimmune disorder in a subject comprising administering to the subject an antibody or antigen-binding fragment thereof that binds to interleukin-22 (IL-22) in an amount sufficient to treat the <u>arthritis</u> autoimmune disorder in the subject.
  - 11. (Cancelled)
- 2. 12. (Currently Amended): The method of claim 10, wherein said <u>arthritis</u> autoimmune disorder is selected from the group consisting of rheumatoid arthritis, osteoarthritis, multiple sclerosis, myasthenia gravis, Crohn's disease, inflammatory bowel disease, lupus-associated arthritis and psoriatic arthritis., diabetes and psoriasis.
- 12 13. (Currently Amended): The method of claim 16, wherein said <u>arthritis</u> autoimmune disorder is rheumatoid arthritis.
- 5 1 2 12 13 14. (Currently Amended): The method of <u>any of claim 10, 12, or 18,</u> wherein said antibody is a neutralizing anti-IL-22 antibody or an antigen-binding fragment thereof.
  - 15. (Cancelled)
- 15. (Original): The method of claim 14, wherein said subject is a human.
- 20 17. (Previously Presented): A method of ameliorating symptoms associated with arthritis, comprising administering to a subject an antibody or antigen-binding fragment thereof that binds to IL-22 in an amount sufficient to ameliorate the symptoms in the subject.

21 18. (Currently Amended): The method of claim 1/2, wherein said arthritis is selected from the group consisting of rheumatoid arthritis, osteoarthritis, lupus-associated arthritis and psoriatic arthritis.

- $1\nu$  19. (Currently Amended): The method of claim 1/2, wherein said IL-22 antibody, or antigen-binding fragment thereof, is administered therapeutically.
- 20 25. (Currently Amended): The method of claim 17, wherein said IL-22 antibody, or antigen-binding fragment thereof, is administered prophylactically.

21-33 (Cancelled)

- 34. (Previously Presented): The method of claim 1/2, wherein said IL-22 comprises an amino acid sequence that is at least 90% identical to amino acids 34-179 of SEQ ID NO:2, wherein said IL-22 is capable of inducing the phosphorylation of a Stat-3 protein.
- 4 36. (Previously Presented): The method of claim 12, wherein said IL-22 comprises an amino acid sequence that is at least 95% identical to amino acids 34-179 of SEQ ID NO:2, wherein said IL-22 is capable of inducing the phosphorylation of a Stat-3 protein.
- 5 %6. (Previously Presented): The method of claim 12, wherein said IL-22 comprises the amino acid sequence of amino acids 34-179 of SEQ ID NO:2.
- 7 3/1. (Previously Presented): The method of claim 1/2, wherein said IL-22 comprises the amino acid sequence of SEQ ID NO:2.
- 8 38. (Previously Presented): The method of claim 12, wherein said antibody, or antigen-binding fragment thereof, binds to a fragment of IL-22 comprising an amino acid sequence selected from the group consisting of amino acids 50-60, 63-81, 84-93, and 168-177 of SEQ ID NO:2.
- 24 39. (Currently Amended): The method of any of claim 17, 18, 19, or 20, wherein said antibody, or antigen-binding fragment thereof, is a neutralizing antibody.
- 9 46. (Previously Presented): The method of claim 12, wherein said antibody, or antigen-binding fragment thereof, is selected from the group consisting of a monoclonal antibody, a polyclonal antibody, a chimeric antibody, a single-chain antibody, a CDR-grafted antibody and a humanized antibody.

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10 41. (Previously Presented): The method of claim 40, wherein said antibody, or antigenbinding fragment thereof, is a monoclonal antibody.

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- 42. (Previously Presented): The method of claim 12, wherein said antibody, or antigen-binding fragment thereof, is a human antibody.
- 4). (Currently Amended): A method of treating rheumatoid arthritis in a subject, comprising administering to the subject an antibody or antigen-binding fragment thereof that binds to IL-22 in an amount sufficient to treat the <u>rheumatoid arthritis</u> autoimmune disorder in the subject, wherein said IL-22 comprises an amino acid sequence that is at least 90% identical to amino acids 34-179 of SEQ ID NO:2 and is capable of inducing the phosphorylation of a Stat-3 protein.
- 42. (Previously Presented): The method of claim 43, wherein said IL-22 comprises an amino acid sequence that is at least 95% identical to amino acids 34-179 of SEQ ID NO:2 and is capable of inducing the phosphorylation of a Stat-3 protein.
- 43 45. (Previously Presented): The method of claim 43, wherein said IL-22 comprises the amino acid sequence of amino acids 34-179 of SEQ ID NO:2.
- 46. (Previously Presented): The method of claim 1/2, wherein said IL-22 comprises an amino acid sequence that is at least 90% identical to amino acids 34-179 of SEQ ID NO:2, wherein said IL-22 is capable of inducing the phosphorylation of a Stat-3 protein.
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  18 47. (Previously Presented): The method of claim 17, wherein said IL-22 comprises an amino acid sequence that is at least 95% identical to amino acids 34-179 of SEQ ID NO:2, wherein said IL-22 is capable of inducing the phosphorylation of a Stat-3 protein.
- 20 29 48. (Previously Presented): The method of claim 17, wherein said IL-22 comprises the amino acid sequence of amino acids 34-179 of SEQ ID NO:2.
- 3/ 46. (Previously Presented): The method of claim 17, wherein said IL-22 comprises the amino acid sequence of SEQ ID NO:2.
- 32 50. (Previously Presented): The method of claim 17, wherein said antibody, or antigen-binding fragment thereof, binds to a fragment of IL-22 comprising an amino acid

sequence selected from the group consisting of amino acids 50-60, 63-81, 84-93, and 168-177 of SEQ ID NO:2.

33 51. (Previously Presented): The method of claim 37, wherein said antibody, or antigen-binding fragment thereof, is selected from the group consisting of a monoclonal antibody, a polyclonal antibody, a chimeric antibody, a single-chain antibody, a CDR-grafted antibody and a humanized antibody.

34 52. (Previously Presented): The method of claim 51, wherein said antibody, or antigenbinding fragment thereof, is a monoclonal antibody.

20 36 58. (Previously Presented): The method of claim 17, wherein said antibody, or antigen-binding fragment thereof, is a human antibody.

36 54. (Previously Presented): The method of claim 17, wherein said arthritis is psoriatic arthritis.

45 56. (Previously Presented): The method of claim 43, wherein said antibody, or antigen-binding fragment thereof, binds to a fragment of IL-22 comprising an amino acid sequence selected from the group consisting of amino acids 50-60, 63-81, 84-93, and 168-177 of SEQ ID NO:2.

4 56. (Previously Presented): The method of claim 43, wherein said antibody, or antigen-binding fragment thereof, is selected from the group consisting of a monoclonal antibody, a polyclonal antibody, a chimeric antibody, a single-chain antibody, a CDR-grafted antibody and a humanized antibody.

47 57. (Previously Presented): The method of claim 56, wherein said antibody, or antigenbinding fragment thereof, is a monoclonal antibody.

48 58. (Previously Presented): The method of claim 43, wherein said antibody, or antigen-binding fragment thereof, is a human antibody.

49 59. (Previously Presented): The method of claim 43, wherein said antibody, or antigen-binding fragment thereof, is a neutralizing antibody.

(Previously Presented): The method of claim 14, wherein said antibody, or antigen-binding fragment thereof, neutralizes IL-22 binding with an ED<sub>50</sub> of about 5nM detected by an enzyme-linked immunoabsorbant assay (ELISA).

25 \$1. (Previously Presented): The method of claim 39, wherein said antibody, or antigen-binding fragment thereof, neutralizes IL-22 binding with an ED<sub>50</sub> of about 5nM detected by an enzyme-linked immunoabsorbant assay (ELISA).

(Previously Presented): The method of claim 59, wherein said antibody, or antigen-binding fragment thereof, neutralizes IL-22 binding with an ED<sub>50</sub> of about 5nM detected by an enzyme-linked immunoabsorbant assay (ELISA).

Please add new claims 63-76:

(New): The method of claim 19, wherein said arthritis is psoriatic arthritis.

17 64. (New): The method of claim 10, wherein said arthritis is osteoarthritis.

18 65. (New): The method of claim 10, wherein said arthritis is associated with lupus.

19 66. (New): The method of claim 10, wherein said antibody or antigen-binding fragment thereof is administered subcutanously or intravenously.

37 67. (New): The method of claim 17, wherein said arthritis is rheumatoid arthritis.

3 % 68. (New): The method of claim 1/7, wherein said arthritis is osteoarthritis.

30 60. (New): The method of claim 17, wherein said arthritis is associated with lupus.

40 76. (New): The method of claim 17, wherein said antibody or antigen-binding fragment thereof is administered subcutanously or intravenously.

7/. (New): The method of claim 30, wherein said subject is a human.

ち フ が. (New): The method of claim 43, wherein said antibody or antigen-binding fragment thereof is administered subcutanously or intravenously.

5) 78. (New): The method of claim 59, wherein said subject is a human.

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(New): The method of claim 36, wherein the subject is a human.

43 44 /5. (New): The method of claim 45, wherein the subject is a human.

76. (New): The method of claim 48, wherein the subject is a human.